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**Wallace et al.**

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[45] **Date of Patent:** **Jan. 11, 2000**

[54] **ZIRCONIUM AND/OR HAFNIUM  
OXYNITRIDE GATE DIELECTRIC**

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provisional application No. 60/053,617, Jul. 24, 1997.

[51] **Int. Cl.**<sup>7</sup> ..... **H01L 21/283**; H01L 21/31;  
H01L 21/441; H01L 21/469

[52] **U.S. Cl.** ..... **438/287**; 438/761; 438/763;  
438/765; 438/769; 438/775; 438/785; 438/786;  
257/213; 257/288; 257/310; 257/410; 257/411

[58] **Field of Search** ..... 438/216, 240,  
438/287, 591, 769, 775, 785, 786; 257/310,  
410, 411

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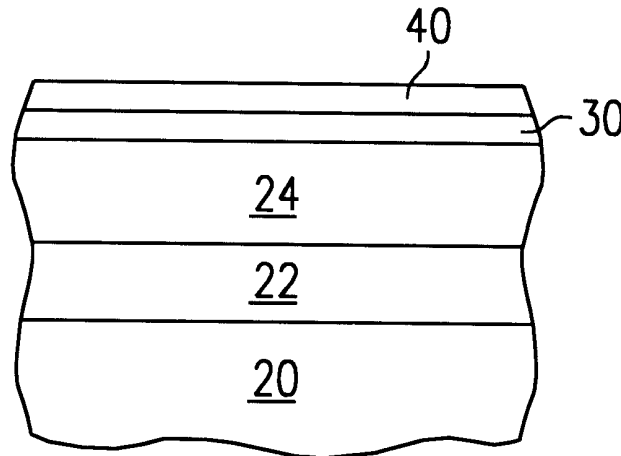
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[57] **ABSTRACT**

A field effect semiconductor device comprising a high  
permittivity zirconium (or hafnium) oxynitride gate dielec-  
tric and a method of forming the same are disclosed herein.  
The device comprises a silicon substrate **20** having a semi-  
conducting channel region **24** formed therein. A zirconium  
oxynitride gate dielectric layer **36** is formed over this  
substrate, followed by a conductive gate **38**. Zirconium  
oxynitride gate dielectric layer **36** has a dielectric constant is  
significantly higher than the dielectric constant of silicon  
dioxide.

**32 Claims, 2 Drawing Sheets**



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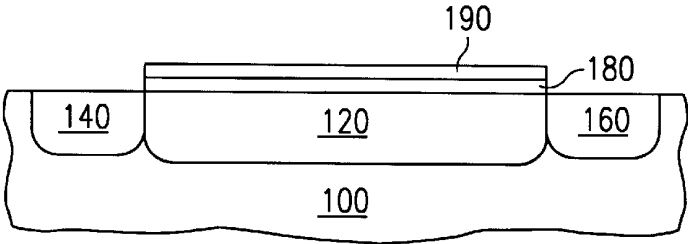


FIG. 1  
(PRIOR ART)

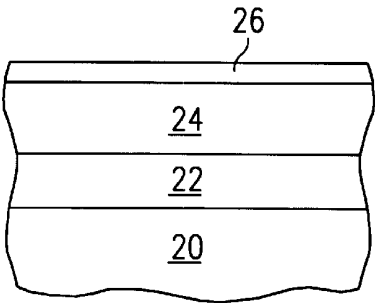


FIG. 2

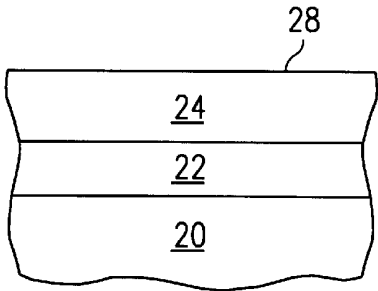


FIG. 3

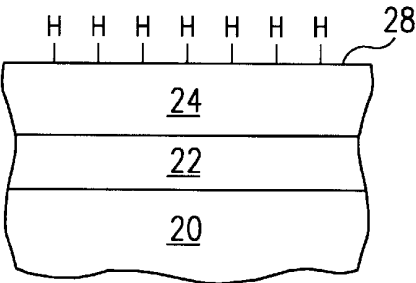


FIG. 4

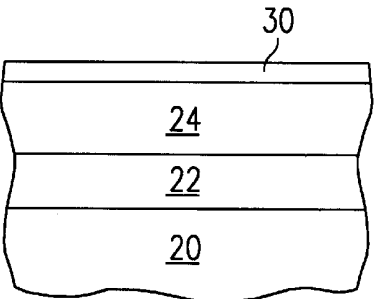


FIG. 5

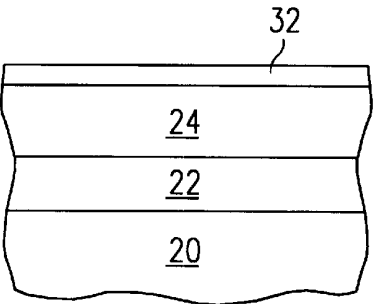


FIG. 6

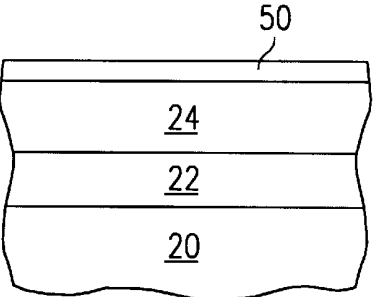


FIG. 7

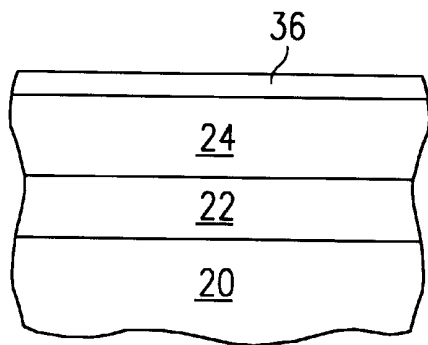


FIG. 8

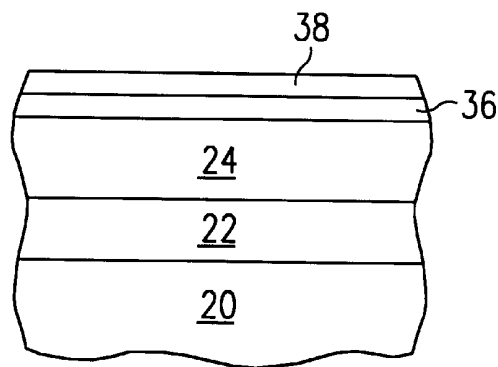


FIG. 9

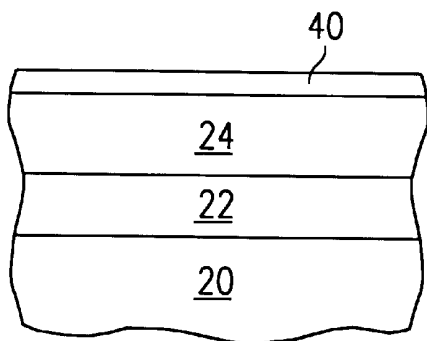


FIG. 10

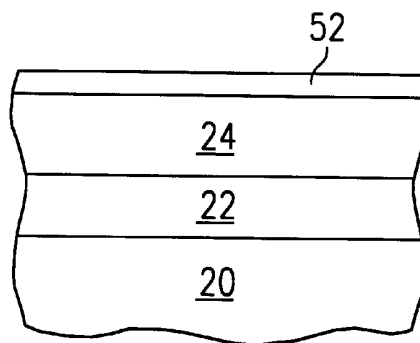


FIG. 11

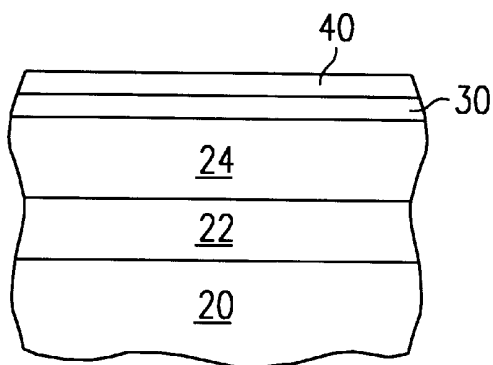


FIG. 12

## ZIRCONIUM AND/OR HAFNIUM OXYNITRIDE GATE DIELECTRIC

This application claims priority from the U.S. provisional application No. 60/053,616, filed on Jul. 24, 1997, now abandoned.

### FIELD OF THE INVENTION

This invention relates generally to semiconductor device structures and methods for forming such, and more specifically to such structures and methods related to gate dielectrics for field effect devices formed on integrated circuits.

### BACKGROUND OF THE INVENTION

Semiconductor devices such as field effect transistors are common in the electronics industry. Such devices may be formed with extremely small dimensions, such that thousands or even millions of these devices may be formed on a single-crystal silicon substrate or "chip" and interconnected to perform useful functions in an integrated circuit such as a microprocessor.

Although transistor design and fabrication is a highly complex undertaking, the general structure and operation of a transistor are fairly simple. With reference to FIG. 1, a simplified field effect transistor is shown in cross-section. In a field effect transistor a portion of the substrate (or epilayer) **100** near the surface is designated as the channel **120** during processing. Channel **120** is electrically connected to source **140** and drain **160**, such that when a voltage difference exists between source **140** and drain **160**, current will tend to flow through channel **120**. The semiconducting characteristics of channel **120** are altered such that its resistivity may be controlled by the voltage applied to gate **190**, a conductive layer overlying channel **120**. Thus by changing the voltage on gate **190**, more or less current can be made to flow through channel **120**. Gate **190** and channel **120** are separated by gate dielectric **180**;

the gate dielectric is insulating, such that between gate **190** and channel **120** little or no current flows during operation (although "tunneling" current is observed with thin dielectrics). However, the gate dielectric allows the gate voltage to induce an electric field in channel **120**, giving rise to the name "field effect transistor."

Generally, integrated circuit performance and density may be enhanced by "scaling", that is by decreasing the size of the individual semiconductor devices on a chip. Unfortunately, field effect semiconductor devices produce an output signal that is proportional to the length of the channel, such that scaling reduces their output. This effect has generally been compensated for by decreasing the thickness of gate dielectric **180**, thus bringing the gate in closer proximity to the channel and enhancing the field effect.

As devices have scaled to smaller and smaller dimensions, the gate dielectric thickness has continued to shrink. Although further scaling of devices is still possible, scaling of the gate dielectric thickness has almost reached its practical limit with the conventional gate dielectric material, silicon dioxide. Further scaling of silicon dioxide gate dielectric thickness will involve a host of problems: extremely thin layers allow for large leakage currents due to direct tunneling through the oxide. Because such layers are formed literally from a few layers of atoms, exacting process control is required to repeatably produce such layers. Uniformity of coverage is also critical because device parameters may change dramatically based on the presence or

absence of even a single monolayer of dielectric material. Finally, such thin layers form poor diffusion barriers to impurities.

Realizing the limitations of silicon dioxide, researchers have searched for alternative dielectric materials which can be formed in a thicker layer than silicon dioxide and yet still produce the same field effect performance. This performance is often expressed as "equivalent oxide thickness": although the alternative material layer may be thick, it has the equivalent effect of a much thinner layer of silicon dioxide (commonly called simply "oxide"). Many, if not most, of the attractive alternatives for achieving low equivalent oxide thicknesses are metal oxides, such as tantalum pentoxide, titanium dioxide, and barium strontium titanate.

Researchers have found formation of such metal oxides as gate dielectrics to be problematic. At typical metal oxide deposition temperatures, the oxygen ambient or oxygen-containing precursor required to form them tends to also oxidize the silicon substrate, producing an oxide layer at the interface between the substrate and the gate dielectric. The presence of this interfacial oxide layer increases the effective oxide thickness, reducing the effectiveness of the alternative gate dielectric approach. The existence of the interfacial oxide layer places an ultimate constraint on the performance of an alternative dielectric field effect device.

### SUMMARY OF THE INVENTION

The present invention includes a semiconductor device structure utilizing either a zirconium oxynitride or a hafnium oxynitride gate dielectric layer, and a method for making the same. This method also encompasses gate dielectrics formed from oxynitrides of mixtures of Zr and Hf. With the present invention, a zirconium (or hafnium) oxynitride gate dielectric may be formed with a dielectric constant substantially higher than that of either conventional thermal silicon dioxide or silicon nitride dielectrics. Thus, the metal (Zr or Hf) oxynitride dielectric layer may be made substantially thicker than a conventional gate dielectric with equivalent field effect. Additionally, the presence of nitrogen, in at least a partial thickness of the gate dielectric, helps to prevent the diffusion of boron, such as from a boron-doped polysilicon gate electrode, to the channel region.

Conventional researcher wisdom has been to avoid nitrogen-based compounds for gate dielectrics. Additionally, integrated circuit manufacturing researchers tend to hesitate before investigating the addition of new materials, and especially material types, for mass produced integrated circuits. In spite of this, our investigations suggest that  $\text{HfO}_x\text{N}_y$  and  $\text{ZrO}_x\text{N}_y$  (with relatively small N levels) are stable next to Si, so there will not be a reaction to form  $\text{SiO}_2$  (silicon oxide reactions are at least minimized to the extent that the dielectric properties are not substantially corrupted). Combining this, and the high permittivities achievable, with our research into understanding the silicon/oxynitride interface has allowed us to recognize the usability of zirconium oxynitride and hafnium oxynitride gate dielectrics.

In one embodiment, a graded zirconium (or hafnium) oxynitride layer is formed, such that near the silicon interface the dielectric layer has a large oxynitride component, while the upper portion of the oxynitride layer has a large zirconium component. Such a structure may employ primarily silicon/oxynitride bonding at the silicon interface, with acceptable interface state densities. However, the zirconium and/or hafnium included in the oxynitride layer can significantly increase the dielectric constant of the film. The present invention also provides for amorphous gate

dielectrics, which have dense microstructures and avoid many of the problems associated with grain boundaries in polycrystalline dielectrics.

In one aspect of this invention, a method of fabricating a semiconductor device is disclosed that includes providing a single-crystal silicon substrate, which usually includes structures, such as a channel region; forming a metal oxynitride gate dielectric layer on the substrate, and forming a conductive gate overlying the gate dielectric layer. This metal can be Zr, hafnium, or a mixture of the two. Suitable base layers for the metal oxynitride layer include bare silicon, silicon oxynitride, and other passivated Si layers, such as hydrogen terminated Si. These other passivation schemes may require removal of the passivant before formation of the zirconium oxynitride.

In one zirconium-based approach, the zirconium oxynitride dielectric layer is formed by forming zirconium on the substrate, and annealing the formed metal in an atmosphere including oxygen and nitrogen, thus forming a metal oxynitride layer on the substrate. In some embodiments, the atmosphere includes NO, or a remote nitrogen/oxygen plasma.

In another zirconium-based approach, the zirconium oxynitride dielectric layer is formed by forming zirconium on the substrate, and annealing the formed metal in a non-oxidizing atmosphere including nitrogen, such as atomic nitrogen or NH<sub>3</sub>, and annealing this zirconium nitride layer in oxygen to form a zirconium oxynitride layer.

In another zirconium-based approach, the zirconium (or hafnium) oxynitride dielectric layer is formed by depositing zirconium or hafnium on the substrate in an atmosphere including oxygen and nitrogen, thus forming a metal oxynitride layer on the substrate. In some embodiments, the atmosphere includes NO, or N<sub>2</sub>O.

#### BRIEF DESCRIPTION OF THE DRAWINGS

This invention, including the features and advantages thereof, can be best understood by reference to the following drawings, wherein:

FIG. 1 is a cross-sectional view of a typical prior art integrated circuit field effect transistor.

FIGS. 2–5 are cross-sectional views of several semiconductor devices, illustrating different surfaces appropriate for deposition of a zirconium oxynitride gate dielectric according to the invention.

FIGS. 6–12 are cross-sectional views of a semiconductor device during fabrication according to various embodiments of the invention.

#### DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

This application describes the embodiments in terms of a zirconium oxynitride dielectric. However, due to its chemical similarity, hafnium can be used in place of zirconium in most of these embodiments. Additionally, hafnium-zirconium mixtures can be used in place of either. In some embodiments, these mixtures will be substantially pure Zr or Hf, with only small amounts of Hf or Zr included. In general, both zirconium oxynitride and hafnium oxynitride provide high dielectric constants and good chemical stability. In some embodiments, Zr will be preferred due to its higher availability and lower cost. However, hafnium oxynitride has a somewhat higher dielectric constant and may be more stable than zirconium oxynitride. This slight performance edge may lead to a preference for hafnium oxynitride in some applications.

The preferred embodiments of the invention may be fabricated on a silicon substrate, as described herein. Although it is not required to practice this invention, it is conventional to use a Si(100) substrate. The description of these embodiments begins, as shown in FIG. 2, after formation of an optional epitaxial Si layer 22 on substrate 20 and implantation of an active channel region 24 in epitaxial layer 22 (or substrate 20, if no epi-layer is used). The description assumes that a protective or native silicon oxide region 26 (preferably comprising less than 1 nm of oxide) overlies channel 24 in the region of interest. Such a silicon oxide layer may be formed by heating a clean substrate to 600–700° C. for approximately 30 seconds, in an oxygen ambient of ~10<sup>-3</sup> Torr. Processes for reaching this step in fabrication are all well known in the art, as are various equivalents to which the present invention is applicable.

The particular embodiment employed for forming a zirconium oxynitride gate dielectric will dictate whether silicon oxide region 26 will be either left in place and used in the formation of an interfacial silicon-oxynitride layer, removed such that the dielectric layer may be directly formed on the underlying silicon, or removed and replaced with a passivation layer designed to inhibit interaction of the substrate in the zirconium oxynitride deposition process.

The immediately following description relates to the preparation of the substrate for deposition of oxynitride-forming materials, and is applicable to the specific embodiments presented thereafter. There are two preferred starting surfaces for the invention if silicon oxide region 26 is to be removed. Region 26 may be removed so as to leave either a clean, bare upper surface 28 as shown in FIG. 3, or a hydrogen-terminated surface 28 as shown in FIG. 4. If oxide region 26 is removed, the bare surface is preferred to the hydrogen-terminated surface if chemical reaction of the highly reactive bare Si surface can be prevented, e.g. by processing in ultrahigh (less than ~10<sup>-8</sup> Torr) vacuum until a point in the particular process where exposure to oxygen can be tolerated. Otherwise, the bare Si surface should be terminated with a suitable passivant, such as hydrogen, which inhibits reoxidation, yet may be readily removed at an appropriate point in the process.

The method of oxide removal is not believed to be critical to the practice of the invention, as long as a clean, oxide-free surface 28 can be maintained until an overlying deposition is performed. One preferred method of performing removal of oxide 26 is by exposure to wet HF, for example by dipping the substrate in dilute HF for 30 seconds and rinsing in deionized water. This both removes the native oxide and hydrogen terminates the surface. Another preferred method is by exposure to HF vapor; this provides similar results, but may be used, e.g., in a cluster tool to further prevent reoxidation or contamination of the surface. Either of these approaches may include other appropriate stripping chemicals, with HF or a NH<sub>4</sub>F solution being preferred as a last step to provide termination.

Several other methods produce a non-terminated surface 28, as shown in FIG. 3. One such method with particular applicability to cluster-tool practice is Si flux desorption. It has been found that below 10<sup>-8</sup> Torr and at 780° C., an Si flux of preferably 1.5 Å/sec for about 600 seconds not only removes native oxide, but produces an atomically smooth, stepped surface that may have advantages for ultrathin gate dielectrics. The alternative is simple desorption by heating of the substrate to high temperature in vacuum or in an H<sub>2</sub> ambient. It is believed, however, that the Si-flux method results in a superior surface structure. In any of these methods, if the substrate will not be kept in ultrahigh

vacuum until an overlying deposition is completed, surface **28** may be hydrogen terminated, e.g. by exposure to atomic hydrogen produced by a plasma or hot filament in an H<sub>2</sub> ambient.

Surface **28** may also be passivated with an ultrathin layer, such as a silicon nitride or silicon oxynitride layer, that is not, strictly speaking, an oxide of silicon. Such layers act as a diffusion barrier and provide oxidation resistance to the substrate during formation of the overlying layer. If an oxynitride layer is used, the preferred method of oxynitridation is by exposure to NO. This layer is preferably less than 1 nm thick, and more preferably less than 0.5 nm thick. Although the reasons are not well understood, this ultrathin passivation layer greatly reduces oxidation of the silicon substrate during formation of an oxygen-bearing gate dielectric.

Although the effective dielectric constant of an oxynitride layer is difficult to measure, it is believed to be in the range of 5 to 6. Thicknesses less than 0.5 nm are even more preferred, in order to lessen the effect of the passivation layer's relatively lower dielectric constant on equivalent oxide thickness. Because nitrogen incorporation is believed to be important to the effectiveness of the oxynitride layer as an oxidation barrier, it may be advantageous to post-anneal a NO-formed oxynitride layer in an ambient that provides atomic nitrogen.

Oxynitrides produced by other methods are not believed to provide sufficient oxidation resistance at the required thicknesses to complete some of the gate dielectric structures disclosed herein, and/or require higher process temperatures, and as such are not preferred. For instance, N<sub>2</sub>O processes result in a much smaller incorporation of N than NO processes. NH<sub>3</sub> processes require a pre-existing SiO<sub>2</sub> film, and thus a uniform sub-nanometer oxynitride film appears to be difficult to achieve using NH<sub>3</sub>. Additionally, NH<sub>3</sub> annealing apparently incorporates undesirable hydrogen into the film structure.

A typical NO process applicable to the present invention is as follows. The substrate is cleaned to remove the pad oxide. As a final step in the cleaning, the substrate is dipped in dilute HF for 30 seconds, and rinsed in deionized water. The substrate is then placed in a reaction chamber, which is then evacuated to 3×10<sup>-8</sup> Torr, and then the substrate is heated to 500° C. to remove the hydrogen passivation from the substrate surface. The substrate is heated to 700° C., and NO at 4 Torr is introduced into the chamber for 10 seconds to form the oxynitride passivation layer. FIG. 5 depicts a passivation layer **30**, e.g., typically an oxynitride, but possibly a nitride passivation layer.

Once the substrate has been prepared to provide either a clean Si surface, an oxide layer, or a protective barrier layer as described above, a zirconium (or hafnium) oxynitride gate dielectric is formed on the substrate by one of several methods. Several of these methods are described below.

We chose zirconium oxynitride and hafnium oxynitride due to their stability next to silicon and higher permittivity. Based on the silicate and oxide data, we believe that these metal oxynitride's heat of formation is more negative than the heat of formation of silicon dioxide and that their permittivity is higher than either SiO<sub>2</sub> or silicon nitride. This should form a more stable gate structure and avoid the preferential formation of interfacial silicon dioxide. Table 1 lists the heat of formation, permittivity, and bandgap of several materials, with silicon dioxide included for comparison purposes.

TABLE 1

Material	Bulk Permittivity	Bandgap (eV)	Heat of Formation (kcal/g/atom of O)
Y <sub>2</sub> O <sub>3</sub>	12		-152
CaO		7	-152
MgO	9.6	7.8	-144
La <sub>2</sub> O <sub>3</sub>	30		-143
SrC		5.8	-142
Ca <sub>3</sub> SiO <sub>5</sub>			-138
Sc <sub>2</sub> O <sub>3</sub>			-137
Ca <sub>2</sub> SiO <sub>4</sub>			-135
HfO <sub>2</sub>	40		-134
ZrO <sub>2</sub>	25	7.8	-131
CeO <sub>2</sub>	26		-129
Al <sub>2</sub> O <sub>3</sub>	10	8.7	-125
Ba <sub>2</sub> SiO <sub>4</sub>			-124
CaSiO <sub>3</sub>			-123
SrSiO <sub>3</sub>			~-123
Mg <sub>2</sub> SiO <sub>4</sub>			-122
Na <sub>2</sub> SiO <sub>3</sub>			-121
BaSiO <sub>3</sub>			-120
MgSiO <sub>3</sub>			-119
ZrSiO <sub>4</sub>			-115
CeSiO <sub>4</sub>			~-115
Bi <sub>4</sub> Si <sub>3</sub> O <sub>12</sub>	35-75		
TiO <sub>2</sub>	30	3-3.5	-110
SiO <sub>2</sub>	3.9	8.9-9.3	-103
Ta <sub>2</sub> O <sub>5</sub>	26	4.5	-100
ZnO	4.6	3.3	-84
WO <sub>3</sub>	42		-66.9
CuO			-37.6
PdO			-21

The zirconium oxynitride may be formed as either a polycrystalline or an amorphous film. Generally, polycrystalline films will have better dielectric constant.—However, amorphous films generally have higher breakdown performance, form a better diffusion barrier, and have lower interface state densities. Also, with many of the embodiments for formation of a zirconium oxynitride dielectric according to the present invention, formation of an amorphous film may be easier than formation of a polycrystalline film because of the uniform stoichiometry required for a polycrystalline film.

#### Embodiment 1

In one embodiment according to the invention, a zirconium or hafnium oxynitride gate dielectric is formed by depositing Zr or Hf on a clean Si surface, oxynitridating this metal layer, and annealing this structure. In this embodiment, a substrate such as shown in either FIG. 3 or FIG. 4 is used. If surface **28** is hydrogen passivated as shown in FIG. 4, the substrate can be briefly heated to above 500° C. in vacuum or an inert ambient to remove the passivation.

Referring to FIG. 6, a metal layer **32** (e.g. zirconium) is deposited directly on surface **28**, e.g. by sputtering, evaporation, chemical vapor deposition (CVD) or plasma CVD. The figures show that surface **28** is the surface of channel region **24**. However, formation of a metal oxynitride layer on the surface of epitaxial layer **22** or substrate **20** follows the same method. Layers **20** and **24** will be used interchangeably below, except where the context shows a distinction is meant.

Sputter deposition is preferably done with a low-energy plasma system, such as collimated or long-throw sputtering. Low deposition rates (e.g. on the order of a few angstroms per second) may be useful, as the total thickness to be deposited is small and uniformity is desired. For an 8" wafer, deposition may be completed in a system with a base pressure of ~10<sup>-8</sup> Torr, an operating pressure of ~10<sup>-3</sup> Torr, and a separation between the sputter gun and the wafer of 16

inches, and the wafer may be rotated to improve uniformity. Ar is an acceptable sputter gas, and the wafer may be maintained at a temperature of 400–600° C. during deposition.

As an alternative to sputtering, metal layer **32** may be deposited by evaporation from an e-beam source onto a substrate at 500–600° C., with a net deposition rate on the order of tenths of angstroms to a few angstroms per second. The substrate is preferably rotated to improve uniformity.

Other alternative methods include CVD or plasma CVD using appropriate precursors, such as zirconium tetrachloride (or hafnium tetrachloride) and hydrogen gas. Nitrogen-bearing zirconium (or hafnium) precursors, such as a Zr (or Hf) nitrate, would also be acceptable. Again, with these methods low deposition rates and temperatures (600° C. and lower) can be useful, and a downstream plasma type reactor is preferred to a reactor where the plasma is generated at the substrate.

Referring now to FIG. **8**, metal layer **32** is converted to a zirconium (or hafnium) oxynitride layer **36** by oxynitridation. Process control is critical during this step, as under-oxynitridation will result in decreased resistivity and diffusion resistance and over-oxynitridation may result in decreased capacitance for layer **36** (due to oxidation of the underlying silicon). Several processes are available for this step. The metal can be converted into zirconium (or hafnium) oxynitride by direct exposure to an oxygen/nitrogen atmosphere, such as NO or N<sub>2</sub>O. Alternatively, a low temperature remote N<sub>2</sub>/O<sub>2</sub> plasma can also be used.

One alternative to direct conversion in an oxynitride atmosphere is nitridation in a nitrogen ambient, such as a remote plasma of nitrogen (preferred) or NH<sub>3</sub>, followed by oxidation. Many oxygen anneal processes are available for this step, such as a low temperature O<sub>2</sub> anneal with or without ultraviolet exposure, or an activated oxygen anneal such as O<sub>3</sub>, O<sub>3</sub> with ultraviolet exposure, a downstream O<sub>2</sub> plasma, N<sub>2</sub>O, or a low temperature O<sub>2</sub> plasma with a DC-biased substrate. As an example of this last process, a downstream 1500W ECR source operating at 1 mTorr, coupled with ~60V DC and 13.56 MHz or 300 kHz RF applied to the substrate may be used while He backside cooling at 80° C. is also applied to the substrate. Processing time is determined experimentally such that both resistivity and dielectric constant lie within an acceptable range.

Generally, a high temperature anneal of zirconium (or hafnium) oxynitride layer **36** is selected to densify or crystallize the film after low temperature oxynitridation. For example, the substrate may be densified by annealing in Ar for 20 seconds at 750° C. This anneal may be done in either an inert or a reducing environment, with a reducing environment particularly useful where metal layer **32** was deposited by CVD using halogens. If a reducing environment is used, an additional low-temperature post-anneal in oxygen may be used to improve dielectric properties of the zirconium (or hafnium) oxynitride layer **36**.

Finally, with reference to FIG. **9**, conductive gate **38** is deposited over the zirconium oxynitride gate dielectric **36**. Processes for depositing gate **38** are well known in the art; gate **38** may be formed, by way of example, of doped polysilicon, metal, or a conductive metal oxide.

In other approaches, polysilicon gates often require an additional passivation layer between dielectric **36** and gate **38** to prevent reduction of dielectric **36** and oxidation of gate **38** at the interface. Although a passivation layer is allowable, the polysilicon reaction with the underlying zirconium oxynitride tends to form an interfacial zone of zirconium silicon-oxynitride, which will not destroy the dielectric

properties of the layer. Thus, many of the compatibility benefits obtained at the bottom interface are also available at the top interface. Additionally, the nitrogen in the dielectric **36** will greatly reduce boron migration through the dielectric **36** to the substrate **20** when a polysilicon gate is doped.

#### Embodiment 2

In a second embodiment according to the invention, a zirconium (or hafnium) oxynitride gate dielectric is formed by depositing zirconium (or hafnium) on a substrate in an oxygen/nitrogen ambient, followed by annealing. This embodiment preferably utilizes a substrate prepared by one of the methods corresponding to FIG. **2**, **3**, or **4**, and the metal may be deposited by one of the methods described in embodiment 1, with the following differences.

Referring to FIG. **10**, an oxynitrided zirconium layer **40** may be deposited on a clean Si surface by sputtering Zr or Hf, as described above. However, some amount of controlled oxygen and nitrogen activity is used to at least partially oxidize-nitridize layer **40** as zirconium is supplied to the substrate. For example, NO or N<sub>2</sub>O may be introduced near the substrate during sputtering with Ar, with a NO flow rate of about one-tenth that of the Ar flow rate. For a metal deposition rate of 0.1 nanometers per second, the oxynitridation gas is preferably introduced from 0 to 5 seconds after the start of the deposition process. This time delay (if present) will help prevent formation of an interfacial SiO<sub>2</sub> layer.

If the zirconium is introduced by the evaporation method, the oxynitridation gas is preferably added near the substrate. To achieve near complete oxynitridation of the deposited metal, ~5–10 Torr of NO may be used for a metal deposition rate of 0.1 nm/sec. If a CVD method is used, appropriate precursors should provide the necessary oxygen (e.g. zirconium tetrachloride and water), and may provide the nitrogen. For a non-nitrogen-bearing precursor, the nitrogen can be provided by a post-deposition nitridation with a nitrogen source such as a remote nitrogen plasma, NO, N<sub>2</sub>O, or NH<sub>3</sub>.

Although this zirconium oxynitride layer will likely react with the underlying Si (either during deposition or during later manufacturing), the resulting zirconium silicon-oxynitride interfacial layer should not present a significant problem for most applications.

In some applications, it may be desirable to increase the thickness of this interfacial zirconium silicon-oxynitride layer. In this approach, the reactions between the zirconium oxynitride and the underlying layer can produce a graded layer. One variation on this method is shown in FIG. **12**, wherein layer **40** is deposited over a silicon oxynitride layer **30**. In such an embodiment, oxygen/nitrogen activity during the anneal may be reduced, and an interfacial zirconium silicon-oxynitride layer **36** may be formed by “stealing” oxygen, nitrogen, and silicon from layer **30** and/or “giving” Zr to the silicon-oxynitride layer **30**. The grading of the structure may be adjusted by adjusting the relative initial thicknesses of layers **30** and **40**.

An alternative method for forming a thicker zirconium silicon-oxynitride interfacial layer is to start the NO before the metal deposition. In this process, the initial NO starts to form a thin silicon oxynitride layer. Introduction of the metal starts the formation of a zirconium oxynitride layer. After annealing, diffusion causes a zirconium silicon-oxynitride interfacial layer. The thickness and gradient of the interfacial layer can be changed by varying the time before starting the metal deposition.

#### Embodiment 3

In a third embodiment according to the invention, a zirconium (or hafnium) oxynitride dielectric is formed by



depositing a metal oxide on the substrate, followed by nitridation. This approach sometimes works better than the silicide approaches above, since the deposited layer is not in a highly reduced (i.e. oxygen deficient) state.

Oxynitrides formed according to this embodiment may be formed on a substrate prepared according to FIGS. 3, 4, or 5.

Referring to FIG. 7, a zirconium oxide layer 50 may be deposited on a clean Si surface by sputtering a metal oxide, such as  $ZrO_2$  or  $HfO_2$ . As an alternative to sputtering, this metal oxide layer 50 may be deposited by evaporation onto the substrate at 500–600° C., with a net deposition rate on the order of tenths of angstroms to a few angstroms per second. The substrate is preferably rotated to improve uniformity.

Referring now to FIG. 11, partially reduced zirconium oxide layer 50 is converted to zirconium oxynitride layer 52 by nitridation. In this method, we sometimes prefer nitridation with a remote plasma of nitrogen. However, some artisans may prefer exposure to  $NO$ , and may even find acceptable results by nitriding with  $N_2O$  or  $NH_3$ .

Post-anneals in  $O_2$  at about 400–550° for up to about 30 minutes generally increase capacitance while maintaining low leakage current. Anneals at higher temperatures or longer times tend to degrade capacitance. Many oxygen anneal processes are available for this step, such as a low temperature  $O_2$  anneal, with or without ultraviolet exposure, or an activated oxygen anneal such as  $O_3$ ,  $O_3$  with ultraviolet exposure, a downstream  $O_2$  plasma,  $N_2O$ , or a low temperature  $O_2$  plasma with a DC-biased substrate. As an approximate example of this last process, a downstream 1500W ECR source operating at 1 mTorr, coupled with ~60V DC and 13.56 MHz or 300 kHz RF applied to the substrate may be used while He backside cooling at 80° C. is also applied to the substrate. Processing time is determined experimentally such that both resistivity and dielectric constant lie within an acceptable range.

Often, it will be desirable to practice a post-formation anneal as described in the first embodiment.

In the approaches mentioned above, it may sometimes be preferable to slow the reactions between the zirconium (or hafnium) oxynitride and the underlying silicon. In these cases, it can be useful to first form a thin silicon oxynitride passivation layer, as described above, on the silicon surface. The zirconium silicon-oxynitride layer can then be formed on this passivation layer with one of the methods described above.

The present invention is not limited by the specific embodiments described herein. Although a particular substrate and type of device have been described herein for clarity, this invention has application to Si-based devices generally which modify the semiconducting characteristics of an active region using the field effect of an overlying conductive region. Various other combinations of the described steps may be used to produce zirconium (or hafnium) oxynitride gate dielectrics, and such are intended to fall within the scope of this invention.

This disclosure has concentrated on gate dielectrics. However, if the leakage current is suitably low, most gate dielectrics can be used as storage dielectrics in a memory cell. It should be noted, that in general, memory dielectrics are not suitable for use as gate dielectrics. This unsuitability is due to the much more demanding interface characteristics that gate dielectrics usually require. With a zirconium (or hafnium) oxynitride dielectrics, the leakage currents can be reduced to acceptable levels by increasing the oxidation time (i.e., the oxygen content of the metal oxynitride). Although

this increased oxygen content slightly decreases the permittivity of the overall structure, the overall dielectric constant is still significantly above current technologies. It should be noted that the physical vapor deposition (PVD) methods above (sputtering and evaporation) typically do not provide a highly conformal layer. Thus, when using one of these PVD methods above for storage dielectric applications, it may not be practical to use this on enhanced surface area geometries, such as rugged polysilicon. However, the high permittivity of zirconium oxynitride allows artisans to manufacture useful, high energy storage, capacitors, without complex surface area enhancement schemes.

What is claimed is:

1. A method of fabricating a field-effect device on an integrated circuit, the method comprising:

providing a single-crystal silicon substrate;

forming a metal oxynitride gate dielectric layer on the substrate,

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof; and

forming a conductive gate overlying the gate dielectric layer.

2. The method of claim 1, wherein the metal is substantially pure zirconium.

3. The method of claim 1, wherein the substrate comprises a clean Si surface immediately prior to the depositing step.

4. The method of claim 1, wherein the forming a metal oxynitride dielectric layer step comprises:

forming a metal on the substrate,

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof; and

annealing the formed metal in an atmosphere including oxygen and nitrogen, thereby forming a metal oxynitride layer on the substrate.

5. The method of claim 4, where the atmosphere includes  $NO$ .

6. The method of claim 4, where the atmosphere includes a nitrogen/oxygen plasma, the plasma generated remotely from the substrate.

7. The method of claim 4, where the annealing time is limited, such that the oxygen to metal ratio of the metal oxynitride layer is lower near the interface with the silicon substrate than near the interface with the conductive gate.

8. The method of claim 4, where the forming a metal step comprises chemical vapor deposition.

9. The method of claim 8, where the chemical vapor deposition uses a nitrogen-bearing precursor.

10. The method of claim 1, wherein the forming a metal oxynitride dielectric layer step comprises:

forming a metal oxide on the substrate,

where the metal is selected from the group of hafnium oxide, zirconium oxide, and mixtures thereof; and

annealing the formed metal oxide in an atmosphere including nitrogen, thereby forming a metal oxynitride layer on the substrate.

11. The method of claim 10, where the atmosphere includes a nitrogen plasma, the plasma generated remotely from the substrate.

12. The method of claim 10, where the atmosphere includes  $NO$ .

13. The method of claim 1, wherein the forming a metal oxynitride dielectric layer step comprises:

forming a metal on the substrate,

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof;

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annealing the formed metal in a non-oxidizing atmosphere including nitrogen, thereby forming a metal nitride layer on the substrate; and

annealing the metal nitride layer in an oxidizing ambient, thereby forming a metal oxynitride layer on the substrate. 5

**14.** The method of claim **13**, wherein the non-oxidizing atmosphere includes atomic nitrogen.

**15.** The method of claim **13**, wherein the non-oxidizing atmosphere includes ammonia. 10

**16.** The method of claim **13**, where the forming a metal step comprises chemical vapor deposition.

**17.** The method of claim **1**, wherein the forming a metal oxynitride dielectric layer step comprises depositing a metal on the substrate in an atmosphere including oxygen and nitrogen, thereby forming a metal oxynitride layer on the substrate;

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof.

**18.** The method of claim **17**, where depositing a metal comprises sputtering material from a target comprised of the metal onto the substrate. 20

**19.** The method of claim **17**, where depositing a metal comprises evaporating the metal from a source.

**20.** The method of claim **17**, where the metal to oxygen ratio is lower near the interface with the silicon substrate than near the center of the metal oxynitride layer. 25

**21.** The method of claim **17**, where the atmosphere includes NO.

**22.** The method of claim **17**, where the atmosphere includes N<sub>2</sub>O. 30

**23.** The method of claim **17**, further comprising annealing the metal oxynitride layer.

**24.** The method of claim **23**, wherein the annealing step is carried out at a temperature sufficient to crystallize the metal oxynitride layer. 35

**25.** The method of claim **1**, wherein the forming a metal oxynitride dielectric layer step comprises forming a silicon oxynitride layer on the substrate; and

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depositing a metal on the silicon oxynitride layer in an atmosphere including oxygen and nitrogen, thereby forming a metal oxynitride layer on the substrate;

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof.

**26.** The method of claim **25**, where the forming a silicon oxynitride layer includes exposing the substrate to a first atmosphere including oxygen and nitrogen; and

the depositing a metal step starts within 5 seconds of the start of the exposing the substrate step.

**27.** The method of claim **1**, wherein the forming a metal oxynitride dielectric layer step comprises forming a silicon oxynitride layer on the substrate;

forming a metal on the silicon oxynitride layer, 15

where the metal is selected from the group of hafnium, zirconium, and mixtures thereof; and

annealing the formed metal in an atmosphere including oxygen and nitrogen, thereby forming a metal oxynitride layer.

**28.** The method of claim **1**, wherein the forming a metal oxynitride dielectric layer step comprises forming a silicon oxynitride layer on the substrate;

forming a metal on the silicon oxynitride layer, where the metal is selected from the group of hafnium, zirconium, and mixtures thereof; 25

annealing the formed metal in a non-oxidizing atmosphere including nitrogen, thereby forming a metal nitride layer on the silicon oxynitride layer; and

annealing the metal nitride layer in an oxidizing ambient, thereby forming a metal oxynitride layer on the silicon oxynitride layer.

**29.** An integrated circuit made by the method of claim **1**.

**30.** An integrated circuit made by the method of claim **4**.

**31.** An integrated circuit made by the method of claim **10**.

**32.** An integrated circuit made by the method of claim **13**.

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